FDA 510(k) Premarket Notification

K790043 TRACOR, CANNULA, BICEPTS BIPOLAR

CLEARED

Feb 8, 1979
34 days to decision

K790043 · Product code: **HIN** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k790043/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coagulator-cutter, Endoscopic, Bipolar (and Accessories) (HIN)
Date received	Jan 5, 1979
Decision date	Feb 8, 1979
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Neomed Corp.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1979

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k790043/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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